evaluating the effectiveness of the AD action and determining whether additional rulemaking action would be needed.

Comments are specifically invited on the overall regulatory, economic, environmental, and energy aspects of the rule that might suggest a need to modify the rule. All comments submitted will be available, both before and after the closing date for comments, in the Rules Docket for examination by interested persons. A report that summarizes each FAA-public contact concerned with the substance of this AD will be filed in the Rules Docket.

Commenters wishing the FAA to acknowledge receipt of their comments submitted in response to this rule must submit a self-addressed, stamped postcard on which the following statement is made: ''Comments to Docket Number 94–NM–216–AD.'' The postcard will be date stamped and returned to the commenter.

The regulations adopted herein will not have substantial direct effects on the States, on the relationship between the national government and the States, or on the distribution of power and responsibilities among the various levels of government. Therefore, in accordance with Executive Order 12612, it is determined that this final rule does not have sufficient federalism implications to warrant the preparation of a Federalism Assessment.

For the reasons discussed above, I certify that this action (1) is not a ''significant regulatory action'' under Executive Order 12866; (2) is not a ''significant rule'' under DOT Regulatory Policies and Procedures (44 FR 11034, February 26, 1979); and (3) will not have a significant economic impact, positive or negative, on a substantial number of small entities under the criteria of the Regulatory Flexibility Act. A final evaluation has been prepared for this action and it is contained in the Rules Docket. A copy of it may be obtained from the Rules Docket at the location provided under the caption **ADDRESSES**.

List of Subjects in 14 CFR Part 39

Air transportation, Aircraft, Aviation safety, Incorporation by reference, Safety.

Adoption of the Amendment

Accordingly, pursuant to the authority delegated to me by the Administrator, the Federal Aviation Administration amends part 39 of the Federal Aviation Regulations (14 CFR part 39) as follows:

PART 39—AIRWORTHINESS DIRECTIVES

1. The authority citation for part 39 continues to read as follows:

Authority: 49 U.S.C. App. 1354(a), 1421 and 1423; 49 U.S.C. 106(g); and 14 CFR 11.89.

§ 39.13 [Amended]

2. Section 39.13 is amended by adding the following new airworthiness directive:

95–02–11 McDonnell Douglas: Amendment 39–9130. Docket 94–NM–216–AD.

Applicability: Model DC–9–87 (MD–87) series airplanes having factory serial numbers (FSN) 49605 through 49612 inclusive, 49614, 53009 through 53011 inclusive, 53336, 53337, 53340, and 53348; certificated in any category.

Note 1: This AD applies to each airplane identified in the preceding applicability provision, regardless of whether it has been modified, altered, or repaired in the area subject to the requirements of this AD. For airplanes that have been modified, altered, or repaired so that the performance of the requirements of this AD is affected, the owner/operator must use the authority provided in paragraph (b) to request approval from the FAA. This approval may address either no action, if the current configuration eliminates the unsafe condition; or different actions necessary to address the unsafe condition described in this AD. Such a request should include an assessment of the effect of the changed configuration on the unsafe condition addressed by this AD. In no case does the presence of any modification, alteration, or repair remove any airplane from the applicability of this AD.

Compliance: Required as indicated, unless accomplished previously.

To prevent severe damage to the airframe in the event of a fire, accomplish the following:

(a) Within 12 months after the effective date of this AD, perform a visual inspection to detect chafing or arcing damage to the wiring of the aft right coatroom, the intercostal, and the recirculation duct assembly near longeron 5 (between stations Y=1078.000 and Y=1098.000), in accordance with McDonnell Douglas MD–80 Service Bulletin 24–151, dated September 29, 1994.

(1) If no damage is found, prior to further flight, modify the wiring installation for the aft right coatroom (reference paragraph 1.C., Condition I, of the service bulletin) in accordance with the procedures described in the service bulletin.

(2) If any damage is found, prior to further flight, modify the wiring installation for the aft right coatroom (reference paragraph 1.C., Condition II, of the service bulletin) in accordance with the procedures described in the service bulletin.

(b) An alternative method of compliance or adjustment of the compliance time that provides an acceptable level of safety may be used if approved by the Manager, Los Angeles ACO. Operators shall submit their requests through an appropriate FAA Principal Maintenance Inspector, who may add comments and then send it to the Manager, Los Angeles ACO.

Note 2: Information concerning the existence of approved alternative methods of compliance with this AD, if any, may be obtained from the Los Angeles ACO.

(c) Special flight permits may be issued in accordance with §§ 21.197 and 21.199 of the Federal Aviation Regulations (14 CFR 21.197 and 21.199) to operate the airplane to a location where the requirements of this AD can be accomplished.

(d) The inspection and modification shall be done in accordance with McDonnell Douglas MD–80 Serivce Bulletin 24–151, dated September 29, 1994. This incorporation by reference was approved by the Director of the Federal Register in accordance with 5 U.S.C. 552(a) and 1 CFR part 51. Copies may be obtained from McDonnell Douglas Corporation, P.O. Box 1771, Long Beach, California 90801–1771, Attention: Business Unit Manager, Technical Administrative Support, Dept. L51, M.C. 2–98. Copies may be inspected at the FAA, Transport Airplane Directorate, 1601 Lind Avenue, SW., Renton, Washington; or at the FAA, Transport Airplane Directorate, Los Angeles Aircraft Certification Office, 3960 Paramount Boulevard, Lakewood, California; or at the Office of the Federal Register, 800 North Capitol Street NW., suite 700, Washington, DC.

(e) This amendment becomes effective on March 2, 1995.

Issued in Renton, Washington, on January 24, 1995.

Darrell M. Pederson,

Acting Manager, Transport Airplane Directorate, Aircraft Certification Service.

[FR Doc. 95–2177 Filed 2–14–95; 8:45 am]

BILLING CODE 4910–13–U

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 178

[Docket No. 91F–0271]

Indirect Food Additives: Adjuvants, Production Aids, and Sanitizers

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the food additive regulations to provide for the safe use of β, 3(or 4)-bis(octadecylthio)cyclohexylethane as an antioxidant for general use in polymeric food-contact articles. This action is in response to a petition filed by Atochem North America, Inc.

DATES: Effective February 15, 1995; written objections and requests for a hearing by March 17, 1995.

ADDRESSES: Submit written objections to the Dockets Management Branch (HFA–305), Food and Drug Administration, rm. 1–23, 12420 Parklawn Dr., Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: Vir Anand, Center for Food Safety and Applied Nutrition (HFS–216), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–254–9500.

SUPPLEMENTARY INFORMATION: In a notice published in the **Federal Register** of August 8, 1991 (56 FR 37712), FDA announced that a food additive petition (FAP 1B4274) had been filed by Atochem North America, Inc., c/o 1150 17th St. NW., Washington, DC 20036 (presently, 1001 G St. NW, suite 500 West, Washington, DC 20001). The petition proposed that the food additive regulations be amended in § 176.170 *Components of paper and paperboard in contact with aqueous and fatty foods* (21 CFR 176.170) to provide for the safe use of β, 3(or 4)-bis(octadecylthio)cyclohexylethane as an antioxidant in polymeric articles intended for food-contact uses. The agency reviewed the nomenclature of the additive and has determined that to ensure unambiguous identification of the compound, a synonym compatible with the Chemical Abstract Service nomenclature, namely, 1-[(*beta*-(octadecylthio)ethyl]-3(or 4)-(octadecylthio)cyclohexane, should be included in this final rule.

Upon further review of the petition, the agency noted that the petitioner had requested use of the additive for general use in polymers rather than as an additive for paper and paperboard. In a notice published in the **Federal Register** of September 11, 1991 (56 FR 46323), FDA amended the filing notice of August 8, 1991, to state that the petitioner had requested that the food additive regulations be amended in § 178.2010 *Antioxidants and/or stabilizers for polymers* (21 CFR 178.2010) to provide for the use of β, 3(or 4)-bis(octadecylthio)cyclohexylethane as an antioxidant for general use in polymeric food-contact articles.

FDA has evaluated the data in the petition and other relevant material. The agency concludes that the proposed use of the additive is safe and that the regulations in § 178.2010 should be amended as set forth below.

In accordance with § 171.1(h) (21 CFR 171.1(h)), the petition and the documents that FDA considered and relied upon in reaching its decision to approve the petition are available for inspection at the Center for Food Safety and Applied Nutrition by appointment with the information contact person listed above. As provided in 21 CFR 171.1(h), the agency will delete from the documents any materials that are not available for public disclosure before making the documents available for inspection.

The agency has carefully considered the potential environmental effects of this action. FDA has concluded that the action will not have a significant impact on the human environment, and that an environmental impact statement is not required. The agency's finding of no significant impact and the evidence supporting that finding, contained in an environmental assessment, may be seen in the Dockets Management Branch (address above) between 9 a.m. and 4 p.m., Monday through Friday.

Any person who will be adversely affected by this regulation may at any time on or before March 17, 1995, file with the Dockets Management Branch (address above) written objections thereto. Each objection shall be separately numbered, and each numbered objection shall specify with particularity the provisions of the regulation to which objection is made and the grounds for the objection. Each numbered objection on which a hearing is requested shall specifically so state. Failure to request a hearing for any particular objection shall constitute a waiver of the right to a hearing on that objection. Each numbered objection for which a hearing is requested shall include a detailed description and analysis of the specific factual information intended to be presented in support of the objection in the event that a hearing is held. Failure to include such a description and analysis for any particular objection shall constitute a waiver of the right to a hearing on the objection. Three copies of all documents shall be submitted and shall be identified with the docket number found in brackets in the heading of this document. Any objections received in response to the regulation may be seen in the Dockets Management Branch between 9 a.m. and 4 p.m., Monday through Friday.

List of Subjects in 21 CFR Part 178

Food additives, Food packaging.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs and redelegated to the Director, Center for Food Safety and Applied Nutrition, 21 CFR part 178 is amended as follows:

PART 178—INDIRECT FOOD ADDITIVES: ADJUVANTS, PRODUCTION AIDS, AND SANITIZERS

1. The authority citation for 21 CFR part 178 continues to read as follows:

Authority: Secs. 201, 402, 409, 721 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321, 342, 348, 379e).

2. Section 178.2010 is amended in the table in paragraph (b) by alphabetically adding a new entry under the headings ''Substances'' and ''Limitations'' to read as follows:

§ 178.2010 Antioxidants and/or stabilizers for polymers.

* * * * *

(b) * * *

Substances	Limitations
* * *	* * * *
β, 3(or 4)-Bis(octadecylthio)cyclohexylethane (CAS Reg. No. 37625–75–5); CAS synonym: 1-[(*beta*-(octadecylthio)ethyl]-3(or 4)-(octadecylthio)cyclohexane.	For use only: 1. At levels not to exceed 0.3 percent by weight of all polymers for use in contact with foods of Types I, II, IV-B, VI, VII-B, and VIII under conditions of use B through H as described in Tables 1 and 2 of § 176.170(c) of this chapter. 2. At levels not to exceed 0.3 percent by weight of polyolefins complying with § 177.1520 of this chapter, for use in contact with food of types III, IV-A, V, VII-A, and IX under conditions of use C through G as described in Tables 1 and 2 of § 176.170(c) of this chapter.
* * *	* * * *

Dated: February 3, 1995.

Fred R. Shank,

Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 95–3804 Filed 2–14–95; 8:45 am]

BILLING CODE 4160–01–F

21 CFR Part 558

New Animal Drugs for Use in Animal Feeds; Tylosin and Virginiamycin

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to remove those portions reflecting approval of four new animal drug applications (NADA's) held by Premiere Agri Technologies, Inc. The NADA's provide for use of Type A medicated articles and Type B medicated feeds containing tylosin and Type B medicated feeds containing virginiamycin. In a notice published elsewhere in this issue of the **Federal Register**, FDA is withdrawing approval of the NADA's.

EFFECTIVE DATE: February 27, 1995.

FOR FURTHER INFORMATION CONTACT: Mohammad I. Sharar, Center for Veterinary Medicine (HFV–216), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–594–1722.

SUPPLEMENTARY INFORMATION: In a notice published elsewhere in this issue of the **Federal Register**, FDA is withdrawing approval of the following NADA's:

NADA No.	Drug name	Sponsor name and address
45–690	Tylosin Type B medicated feeds and Type A medicated article.	Premiere Agri Technologies, Inc., P.O. Box 2508, Fort Wayne, IN 46801–2508 (former sponsor Henwood Feed Additives)
97–289	Tylosin Type B medicated feeds and Type A medicated article.	Do. (Former sponsor Feed Specialties Co., Inc.)
133–361	Virginiamycin Type B medicated feed.	Do. (Former sponsor Feed Specialties Co., Inc.)
133–839	Virginiamycin Type B medicated feed.	Do. (Former sponsor MacPage, Inc.)

The sponsor requested withdrawal of approval of the NADA's. This final rule removes 21 CFR 558.625(b)(11) and (b)(15) and amends 21 CFR 558.635(b)(2) to reflect the withdrawal of approval of these NADA's.

List of Subjects in 21 CFR Part 558

Animal drugs, Animal feeds.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs and redelegated to the Center for Veterinary Medicine, 21 CFR part 558 is amended as follows:

PART 558—NEW ANIMAL DRUGS FOR USE IN ANIMAL FEEDS

1. The authority citation for 21 CFR part 558 continues to read as follows:

Authority: Secs. 512, 701 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b, 371).

§ 558.625 [Amended]

2. Section 558.625 *Tylosin* is amended by removing and reserving paragraphs (b)(11) and (b)(15).

3. Section 558.635 *Virginiamycin* is amended by revising paragraph (b)(2) to read as follows:

§ 558.635 Virginiamycin

* * * * *

(b) * * *

(2) 2.2 percent activity (10 grams per pound) to 011490, 016968, and 017790 in § 510.600(c) of this chapter for use as in paragraphs (f)(1)(iv) and (f)(1)(v) of this section.

* * * * *

Dated: January 6, 1995.

Stephen F. Sundlof,

Director, Center for Veterinary Medicine.

[FR Doc. 95–3802 Filed 2–14–95; 8:45 am]

BILLING CODE 4160–01–F

UNITED STATES INFORMATION AGENCY

22 CFR Part 514

[Rulemaking No. 110]

Exchange Visitor Program

AGENCY: United States Information Agency.

ACTION: Final rule.

SUMMARY: The Agency hereby adopts as final with modifications the interim rule governing its oversight and administration of au pair programs. Au pair programs permit foreign nationals to enter the United States for a period of one year for the purpose of residing with an American host family while participating directly in the home life of the family and providing limited child care services. The foreign national also attends a United States accredited post-secondary educational institution. These rules are promulgated pursuant to Public Law 103–415 which authorizes the continued operation, until September 30, 1995, of au pair programs currently designated by the Agency.

DATES: Effective date: These rules are effective February 15, 1995.

Applicability dates: With the exceptions of § 514.31(j) (1) and (4), and § 514.31(k), these rules apply to all au pair placements and operations as of February 15, 1995. The provisions set forth at § 514.31(j) (1) and (4) and § 514.31(k) shall apply only to au pair participants placed after date of publication.

Compliance date: Sponsor implementation of the provisions set forth at § 514.31(g) (1) and (2) will not be expected before March 31, 1995.

FOR FURTHER INFORMATION CONTACT: Stanley S. Colvin, Assistant General Counsel, United States Information Agency, 301 4th Street, SW., Washington, DC 20547; Telephone, (202) 619–6829.

SUPPLEMENTARY INFORMATION: First begun pursuant to the provisions of the United States Information and Educational Exchange Act of 1948 (''Smith-Mundt''), and subsequently incorporated into and broadened under the Fulbright-Hays Act, educational and cultural exchange activities have, over the past forty years, exposed millions of foreign nationals to the United States, its peoples, cultures, skills, business techniques, educational institutions, and way of life. The Fulbright-Hays Act mandates reciprocal exchange and Americans traveling abroad have, in similar fashion, developed an enhanced awareness of foreign people, their cultures and societies. Thus, Fulbright-Hays programs further one of the Agency's primary missions: increasing mutual understanding between Americans and others through people-to-people contact. Originally conducted by the Department of State, oversight of exchange activities, occurring under the umbrella of the Exchange Visitor Program, has been the responsibility of the Agency since 1978.

The Fulbright-Hays Act sets forth certain parameters which all exchange activities must meet. With an eye towards ensuring that these parameters were being met and acting in response to a Congressional request, the General Accounting Office (''GAO'') investigated Agency oversight and administration of the Exchange Visitor Program and its attendant utilization of the J visa. In its report to Congress, dated February 5, 1990 and entitled ''Inappropriate Uses